Under the Paperwork Reduction Act of 1995, no persons are required to re

INFORMATION DISCLOSURE STATEMENT BY APPLICANT (Not for submission under 37 CFR 1.99)

Application Number		10597648			
Filing Date		2006-08-02			
First Named Inventor	Hidea	ki KIKO			
Art Unit		2161			
Examiner Name	Unas	signed			
Attorney Docket Number		AIBARA0003			

					,							
					U.S.	PATENTS			Remove		_	
Examiner Initial*	Cite No	Patent Number	Kind Code ¹				Name of Patentee or Applicant R			Pages,Columns,Lines where Relevant Passages or Relevant Figures Appear		
	1											
If you wisl	h to a	dd additional U.S. Pate	nt citatio	n inform	ation pl	lease click the	Add button.	_	Add		_	
			U.S.P	ATENT	APPLI	CATION PUB	LICATIONS		Remove			
Examiner Initial*				deited Decument Releva		s,Columns,Lines where ant Passages or Relevant es Appear						
	1											
If you wisl	h to a	dd additional U.S. Publi	shed Ap	plication	citatio	n information p	please click the Ad	d button	Add			
				FOREIG	GN PAT	ENT DOCUM	ENTS		Remove			
Examiner Initial*	Cite No	Foreign Document Number ³	Country Code ²		Kind Code ⁴	Publication Date	Name of Patente Applicant of cited Document	e or	where Rel	or Relevant	TE	
	1	1462945	EP		A1	2004-09-29	TERADA, Shinji				×	
	2	03/048945	wo		A1	2003-06-12	CYBIRD CO., LTD				×	
If you wis	h to a	l dd additional Foreign P				information pl		button	Add			
			NON	e-PATE	NI CITE	KATUKE DO	CUMENTS		Collove	J		

INFORMATION DISCLOSURE STATEMENT BY APPLICANT (Not for submission under 37 CFR 1.99)

pplication Number		10597648		
iling Date		2006-08-02		
irst Named Inventor Hidea		kı KIKO		
ut Unit		2161		
xaminer Name Unas		signed		
ttorney Docket Numb	er	AIBARA0003		

Examiner Initials*	Cite No	include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc), date, pages(s), volume-issue number(s), publisher, city and/or country where publisher.	Ţ5				
	1	ATSUYA YCSHIDA. Virtual Chet, XP-002986895, Nikkei Business Publications, September 9, 1998, pages 151-159, no. 670, Tokyo, JAPAN.					
	2	Supplementary European Search Report Issued in the Corresponding Application PCT/JP2004/003866, Completed July 11, 2007.	×				

If you wish to add additional non-patent literature document citation information please click the Add button Add

Examiner Signature Date Considered

F

A

A

*EXAMINER: Initial if reference considered, whether or not citation is in conformance with MPEP 609. Draw line through a citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

1 See Kind Codes of USPTO Patient Documents at year LISPTO_CODE or MPEP 901.6.2 Enter office that issued the document, by the holidate code (WIPO Standard ST.3). Story duplance patient colourests, by mischool on the year of the Proprior many percectible see-sel number of the period recomment, provided the part of the Proprior many percectible see-sel number of the period recomment, "kind of document by the appropriate symbols as indicated on the document under WIPO Standard ST.16 if possible. "Applicant is to place a check mark here if English languages the relation in a standard."

INFORMATION DISCLOSURE STATEMENT BY APPLICANT (Not for submission under 37 CFR 1.99)

Application Number		10597648	
Filing Date		2006-08-02	
First Named Inventor	Hideakı KIKO		
Art Unit		2161	
Examiner Name	Unass	signed	
Attorney Docket Numb	er	AIBARA0003	

CERTIFICATION STATEMENT

Please see	37	CFR .	1 97	and	1 98 to	make	the ·	enntont	ista ca	laction/	e١٠

That each item of information contained in the information disclosure statement was first cited in any communication from a foreign patent office in a counterpart foreign application not more than three months prior to the filling of the information disclosure statement. See 37 CFR 1.97(e/11).

OR

That no item of information contained in the information disclosure statement was cited in a communication from a foreign patent office in a counterpart foreign application, and, to the knowledge of the person signing the certification after making reasonable inquity, no item of information contained in the information disclosure statement was known to any individual designated in 37 CFR 156(c) more than three months prior to the filing of the information disclosure statement. See 37 CFR 157(e).

- See attached certification statement.
- Fee set forth in 37 CFR 1.17 (p) has been submitted herewith.

SIGNATURE

A signature of the applicant or representative is required in accordance with CFR 1.33, 10.18. Please see CFR 1.4(d) for the form of the signature.

Signature	/Joerg-Uwe Szipl/	Date (YYYY-MM-DD)	2007-08-31
Name/Brint	James I burn Carlot	Registration Number	21700

This collection of information is required by 3T CFR 1.97 and 1.98. The information is required to obtain or retain a benefit by the public which is to file fand by the USPTO to process) an application. Confidentially is governed by \$5 U.S. C. 12 and 37 CFR 1.14. This collection is estimated to take 1 hour to complete, including gathering, preparing and submitting the completed application from the USPTO. Time will vary depending upon the individual case. Any comments on the amount of time you require to complete this form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patent and Trademark Office, u.S. Operatment of Commence, P. 0. Box 1445, Alexandris, V.S. 2231-1450. D. NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Commissioner for Patents, P.O. Box 1450, Alexandria, V.S. 2231-1450.

Privacy Act Statement

The Privacy Act of 1974 (P. L. 93-579) requires that you be given certain information in connection with your submission of the stackhold from related to a patient application or patient. Accordingly, pursuant to the requirements of the Act, places be advised that (1) the general authority for the collection of this information is 35 U.S.C. 2(b)(2); (2) familishing of the information solicided is couldrain; and (3) the primoral pursuance for which the information is used by the U.S. Patient and Trademan Coffice is to process and/or examine your submission related to a patient agricultant or patient. If you do not furnish the requested process and/or examine your submission related to a patient agricultant or patient. If you do not furnish the requested results of the patient of the patient and the patient of the patient

The information provided by you in this form will be subject to the following routine uses:

- The information on this form will be treated confidentially to the extent allowed under the Freedom of Information Act (5 U.S.C. 552) and the Privacy Act (5 U.S.C. 552a). Records from this system of records may be disclosed to the Department of Justice to determine whether the Freedom of Information Act requires disclosure of these record s.
 - A record from this system of records may be disclosed, as a routine use, in the course of presenting evidence to a court, magistrate, or administrative tribunal, including disclosures to opposing counsel in the course of settlement negotiation.
 - A record in this system of records may be disclosed, as a routine use, to a Member of Congress submitting a request involving an individual, to whom the record pertains, when the individual has requested assistance from the Member with respect to the subject matter of the record.
 - A record in this system of records may be disclosed, as a routine use, to a contractor of the Agency having need for the information in order to perform a contract. Recipients of information shall be required to comply with the requirements of the Privacy Act of 1974, as amended, pursuant to 5 U.S.C. 552(m).
 - A record related to an International Application filed under the Patent Cooperation Treaty in this system of records
 may be disclosed, as a routine use, to the International Bureau of the World Intellectual Property Organization, pursuant
 to the Patent Cooperation Treaty.
 - A record in this system of records may be disclosed, as a routine use, to another federal agency for purposes of National Security review (35 U.S.C. 181) and for review pursuant to the Atomic Energy Act (42 U.S.C. 218(c)).
 - 7. A record from this system of records may be disclosed, as a routine use, to the Administrator, General Services, or hisher designed, uturing an insection of records conducted by GSA as part of that agency's responsibility to recommend improvements in records management practices and programs, under authority of 4d U.S.C. 2904 and 2905. Such disclosure shall be made in accordance with the GSA requisions governing inseption of records for this purpose, and any other relevant (i.e., GSA or Commerce) directive. Such disclosure shall not be used to make determinations about individuals.
- A record from this system of records may be disclosed, as a routine use, to the public after either publication of the
 application pursuant to 35 U.S.C. 12(2) to rissuance of a patent pursuant to 35 U.S.C. 151. Further, a record may be
 disclosed, subject to the limitations of 37 CFR 1.14, as a routine use, to the public if the record was filled in application
 which became abandoned or in which the proceedings were terminated and which application is referenced by either a
 published application, an application open to public inspections or as issued patent.
 - A record from this system of records may be disclosed, as a routine use, to a Federal, State, or local law enforcement agency, if the USPTO becomes aware of a violation or potential violation of law or regulation.